WHAT IS CLAIMED IS:

1	1. A method of treating a subject having a pathologic condition
2	involving neovascularization comprising administering a pharmaceutical preparation
3	comprising an R'-Glu-Trp-R" dipeptide and a pharmaceutically acceptable carrier to the
4	subject in an amount effective to inhibit neovascularization.
1	2. The method of claim 1 wherein the R'-Glu-Trp-R" dipeptide is L-
2	Glu-L-Trp.
1	3. The method of claim 2 wherein the condition is hemangioma.
1	4. The method of claim 2 wherein the condition is vascularized
2	malignant tumor or vascularized benign tumor.
1	5. The method of claim 2 wherein the condition is neovascularization
2	in post-recovery cerebrovascular accident; neovascularization due to head trauma;
3	restenosis following angioplasty; or negvascularization due to heat or cold trauma.
1	6. The method of claim 2 wherein the condition is neovascularization
2	associated with substance-induced neovascularization of the liver, angiogenic
3	dysfunction related to an excess of hormone; neovascular sequelae of diabetes;
4	neovascular sequelae to hypertension; or chronic liver infection.
1	7. The method of claim 2 comprising administering to the subject a
2	dose of about 0.5 ag per 1 kilogram body weight to about 1 mg per 1 kg body weight.
1	8. The method of claim 7 wherein the effective amount is about
2	1 μ g/kg to/about 50 μ g/kg body weight.
1	9. The method of claim 7 wherein the dose is administered daily
2	over a period of 1 day to about 30 days.

1	10. The method of claim 7 wherein the pharmaceutical preparation is
2	administered intramuscularly or intranasally.
1	11. The method of claim 2 comprising administering the
2	pharmaceutical preparation in the form of an injectable solution containing 0.001% to
3	0.01% of L-Glu-L-Trp.
1	12. The method of claim 2 comprising administering the preparation
2	in a unit dose form comprising a tablet, a suppository, a capsule, an eye film, an
3	inhalant, a mucosal spray, a nose drop, an eye drop, a toothpaste, an ointment, or water
4	soluble based cream.
1	13. The method of claim 12 wherein said unit dose form consists
2	essentially of 0.01 mg of said R'-Glu-Tfp-R".
1	14. The method of claim 2 further comprising administering to the
2	subject a vasoactive drug.
1	15. The method of claim 14 wherein the vasoactive drug is an
2	angiotensin converting enzyme (ACE) inhibitor or a potassium channel opener (PCO).
1	76. The method of claim 2 wherein the subject suffers from a tumor
2	wherein the method further comprises administering a chemotherapeutic agent.
l	17. The method of claim 2 wherein the subject is not immune
2	compromised.
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